

Product Datasheet

JUNB (Phospho-S259) Rabbit Polyclonal Antibody (orb256641)

Catalog Number	orb256641
Category	Antibodies
Description	The JUNB (Phospho-S259) Antibody is suitable for IHC, IP, WB. It is a Polyclonal, Unconjugated antibody which raised against KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S259 of human JUNB protein. The exact sequence is proprietary. Purification: The antibody was purified by immunogen affinity chromatography.
Target	JUNB
Clonality	Polyclonal
Species/Host	Rabbit
Conjugation	Unconjugated
Reactivity	Bovine, Human, Mouse, Rat, Sheep
Form/Appearance	Liquid
Buffer/Preservatives	0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.
Purification	The antibody was purified by immunogen affinity chromatography.
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S259 of human JUNB protein. The exact sequence is proprietary.
UniProt ID	P09450, P17275, P24898
Tested applications	IHC, IP, WB

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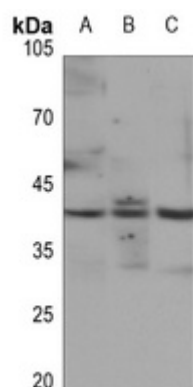
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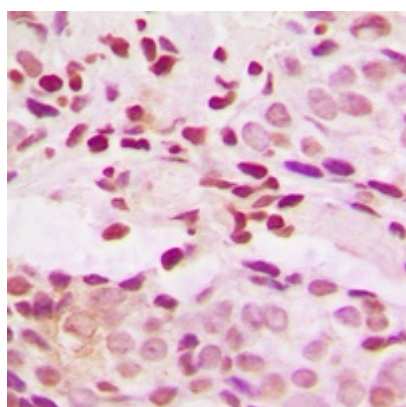
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Dilution range	WB: 1:500-1000, IHC-P: 1:100-200, IP: 1:10-100
Specificity	Recognizes endogenous levels of JUNB protein only when phosphorylated at S259.
Antibody Type	Primary Antibody
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Note	For research use only
Entrez	16477, 24517, 3726
Expiration Date	12 months from date of receipt.



Western blot analysis of JUNB (Phospho-S259) expression in mouse liver (A), mouse kidney (B), rat kidney (C) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 43 kD)



Immunohistochemical analysis of JUNB (Phospho-S259) 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 (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.

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