

## Product Datasheet

### SAF-B1 Antibody (orb412667)

<b>Catalog Number</b>	orb412667
<b>Category</b>	Antibodies
<b>Description</b>	The SAF-B1 Antibody is suitable for IF, IHC, WB. It is a Polyclonal, Unconjugated antibody which raised against KLH-conjugated synthetic peptide of human SAF-B1 .Purification: The antibody was purified by immunogen affinity chromatography.
<b>Clonality</b>	Polyclonal
<b>Species/Host</b>	Rabbit
<b>Conjugation</b>	Unconjugated
<b>Reactivity</b>	Human, Mouse, Rat
<b>Form/Appearance</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.
<b>Purification</b>	The antibody was purified by immunogen affinity chromatography.
<b>Immunogen</b>	KLH-conjugated synthetic peptide of human SAF-B1
<b>UniProt ID</b>	<b>Q15424, O88453, D3YXK2</b>
<b>Tested applications</b>	IF, IHC, WB
<b>Dilution range</b>	WB: 1:500-2000
<b>Specificity</b>	Recognizes endogenous levels of SAF-B1 protein.
<b>Antibody Type</b>	Primary Antibody

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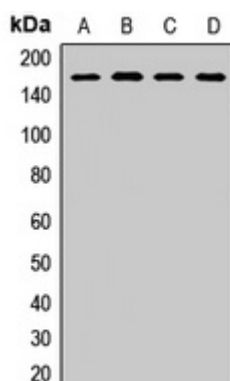
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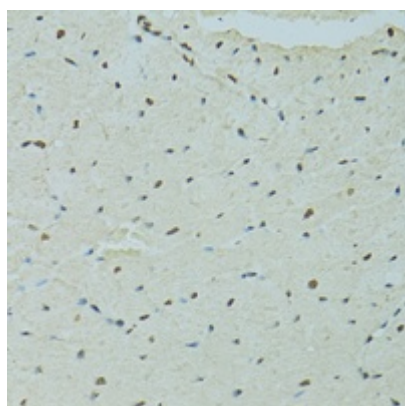
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<b>Storage</b>	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
<b>Note</b>	For research use only
<b>Entrez</b>	<b>224903, 6294, 64196</b>
<b>Expiration Date</b>	12 months from date of receipt.



Western blot analysis of SAF-B1 expression in Hela (A), NIH3T3 (B), mouse lung (C), rat brain (D) whole cell lysates. (Predicted band size: 95; 102 kD; Observed band size: 155 kD)



Immunohistochemical analysis of SAF-B1 staining in mouse 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.

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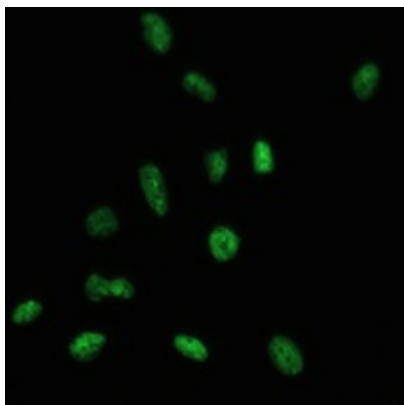
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Immunofluorescent analysis of SAF-B1 staining in U2OS cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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