

Product Data Sheet

Anti-KSR1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4449	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Rat	obit polyclonal antibody	y to KSR1		
Immunogen	KLł	I-conjugated synthetic	peptide encompassing a sequence within the center		
	reg	ion of human KSR1. The	e exact sequence is proprietary.		
Purification	The	e antibody was purified	by immunogen affinity chromatography.		
Specificity	Red	cognizes endogenous le	vels of KSR1 protein.		
Clonality	Pol	yclonal			
Form	Liq	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	d 0.01% sodium azide.			
Dilution	WE	3 (1/500 - 1/1000), IH (1/:	100 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	KSF	₹1			
Alternative N	ames KSF	KSR; Kinase suppressor of Ras 1			
Entrez Gene	884	8844 (Human); 16706 (Mouse)			
SwissProt	Q8	IVT5 (Human); Q61097	(Mouse)		
Storage/Stabi	lity Shi	pped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	fre	eze/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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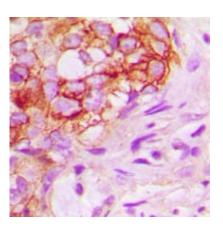
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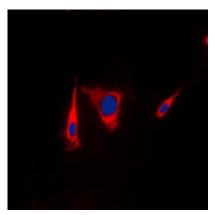
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Western blot analysis of KSR1 expression in HEK293T (A), NIH3T3 (B) whole cell lysates.



Immunohistochemical analysis of KSR1 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of KSR1 staining in HEK293T 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 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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