

Product Data Sheet

Anti-EIF2S1 Antibody

Catalog #	Source	Reactivity	Applications		
-		-			
CPA1376	Rabbit	H, M, R, B, C, P, Z	WB, IH		
Description	Ra	abbit polyclonal antibody to	EIF2S1		
Immunogen	KL	H-conjugated synthetic pep	tide encompassing a sequence within the center		
	re	gion of human EIF2S1. The e	exact sequence is proprietary.		
Purification	Th	ne antibody was purified by i	mmunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous levels	of EIF2S1 protein.		
Clonality	Ро	blyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/50 -	1/100)		
Gene Symbol	EII	F2S1			
Alternative Na	ames Ell	F2A; Eukaryotic translation i	nitiation factor 2 subunit 1; Eukaryotic translation		
	ini	itiation factor 2 subunit alph	a; eIF-2-alpha; eIF-2A; eIF-2alpha		
Entrez Gene	19	965 (Human); 13665 (Mouse); 54318 (Rat)		
SwissProt	PC)5198 (Human); Q6ZWX6 (M	ouse); P68101 (Rat)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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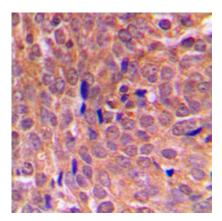
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Western blot analysis of EIF2S1 expression in HEK293T (A), SGC7901 (B), rat heart (C), mouse heart (D) whole cell lysates. (Predicted band size: 36 kD; Observed band size: 38 kD)



Immunohistochemical analysis of EIF2S1 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.

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