

# **Product Data Sheet**

## **Anti-ArgRS Antibody**

Catalog #	Source	Reactivity	Applications		
CQA1331	Rabbit	H, M, R	WB, IH, IF/IC		
Description	I	Rabbit polyclonal antibody	to ArgRS		
Immunogen	I	Recombinant full length pro	otein of human ArgRS		
Purification	-	The antibody was purified b	by immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of ArgRS protein.			
Clonality	I	Polyclonal			
Conjugation					
Form	I	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/2000), IH (1/50	) - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	I	RARS			
Alternative N	ames /	ArgininetRNA ligase, cytop	plasmic; Arginyl-tRNA synthetase; ArgRS		
Entrez Gene	ļ	5917 (Human); 104458 (Mc	ouse); 287191 (Rat)		
SwissProt	I	254136 (Human); Q9D0I9 (I	Mouse); P40329 (Rat)		
Storage/Stabi	i <b>lity</b> S	Shipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	f	reeze/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

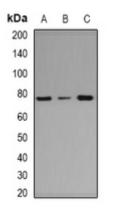
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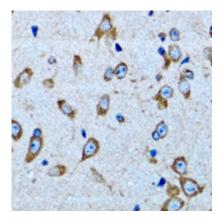


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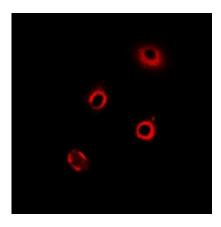
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Western blot analysis of ArgRS expression in Hela (A), HepG2 (B), mouse spleen (C) whole cell lysates. (Predicted band size: 67; 75 kD; Observed band size: 75 kD)



Immunohistochemical analysis of ArgRS staining in rat 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.



Immunofluorescent analysis of ArgRS staining in MCF7 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.

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