

Product Data Sheet

Anti-IARS2 Antibody

Catalog #	Source	Reactivity	Applications	
CPA2714	Rabbit	H, M, R, Mk	WB, IH, IF/IC	
Description	R	abbit polyclonal antibody to	DIARS2	
Immunogen	K	LH-conjugated synthetic pe	otide encompassing a sequence within the N-term	
	re	region of human IARS2. The exact sequence is proprietary.		
Purification	TI	he antibody was purified by	immunogen affinity chromatography.	
Specificity	R	ecognizes endogenous leve	ls of IARS2 protein.	
Clonality	Po	olyclonal		
Conjugation				
Form	Li	iquid in 0.42% Potassium pł	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
	aı	nd 0.01% sodium azide.		
Dilution	W	VB (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)	
Gene Symbol	IA	ARS2		
Alternative Na	a <mark>mes</mark> Is	oleucinetRNA ligase mito	chondrial; Isoleucyl-tRNA synthetase; IleRS	
Entrez Gene	5!	5699 (Human); 381314 (Mc	use)	
SwissProt	Q	9NSE4 (Human); Q8BIJ6 (N	ouse)	
Storage/Stabi	lity Sł	hipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
	fr	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

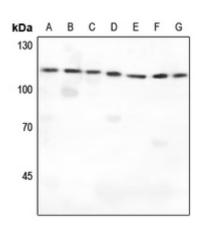
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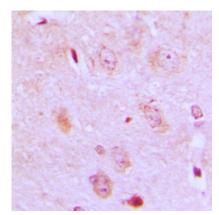
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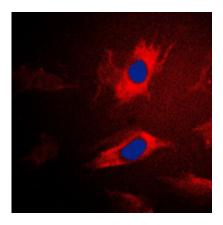
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Western blot analysis of IARS2 expression in HEK293T (A), Hela (B), H446 (C), mouse kidney (D), mouse testis (E), rat kidney (F), rat testis (G) whole cell lysates. (Predicted band size: 113 kD; Observed band size: 114 kD)



Immunohistochemical analysis of IARS2 staining in human 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 IARS2 staining in Jurkat 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 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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