

Product Data Sheet

Anti-ARRDC3 Antibody

Catalog #	Source	Reactivity	Applications
CPA2736	Rabbit	H, M, R, Mk	WB, IH, IF/IC, IP
Description	Ra	bbit polyclonal antibody to	ARRDC3
Immunogen	KLI	H-conjugated synthetic pe	otide encompassing a sequence within the C-term
	reg	gion of human ARRDC3. Th	e exact sequence is proprietary.
Purification	Th	e antibody was purified by	immunogen affinity chromatography.
Specificity	Re	cognizes endogenous leve	s of ARRDC3 protein.
Clonality	Ро	lyclonal	
Form	Liq	uid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	d 0.01% sodium azide.	
Dilution	W	B (1/500 - 1/1000), IH (1/100) - 1/200), IF/IC (1/100 - 1/500), IP (1/10 - 1/100)
Gene Symbol	AR	RDC3	
Alternative N	ames KIA	AA1376; Arrestin domain-c	ontaining protein 3; TBP-2-like inducible membrane
	pro	otein; TLIMP	
Entrez Gene	57	561 (Human); 105171 (Mo	use); 309945 (Rat)
SwissProt	Q9	96B67 (Human); Q7TPQ9 (1	Mouse); Q6TXF1 (Rat)
Storage/Stabi	i lity Shi	ipped at 4°C. Upon deliver	y 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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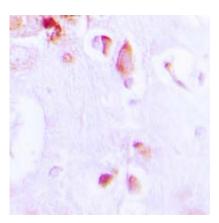
A

B C

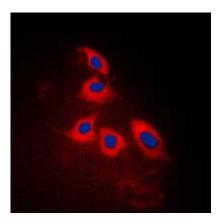
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Western blot analysis of ARRDC3 expression in Jurkat (A), mouse liver (B), rat liver (C) whole cell lysates.



Immunohistochemical analysis of ARRDC3 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 ARRDC3 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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