

Product Data Sheet

Anti-GLURD2 Antibody

Catalog #	Source	Reactivity	Applications
CPA3976	Rabbit	H, M, R, Z	WB, IF/IC
Description	Rabl	oit polyclonal antibody t	o GLURD2
Immunogen	KLH-	conjugated synthetic pe	ptide encompassing a sequence within the C-term
	regio	on of human GLURD2. Th	ne exact sequence is proprietary.
Purification	The	antibody was purified by	<i>immunogen affinity chromatography.</i>
Specificity	Reco	ognizes endogenous leve	ls of GLURD2 protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/1000), IF/IC (1/	100 - 1/500)
Gene Symbol	GRID	02	
Alternative Na	ames GLU	RD2; Glutamate recepto	r ionotropic, delta-2; GluD2; GluR delta-2 subunit
Entrez Gene	2895	5 (Human); 14804 (Mous	se); 79220 (Rat)
SwissProt	O43	424 (Human); Q61625 (N	Mouse); Q63226 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid
	free	ze/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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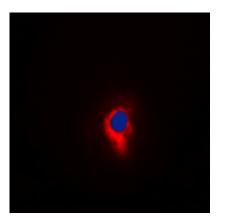
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35 25 For research purposes only, not for human use

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Western blot analysis of GLURD2 expression in zebrafish (A) whole cell lysates. (Predicted band size: 113 kD; Observed band size: 113 kD)



Immunofluorescent analysis of GLURD2 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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