

## **Product Data Sheet**

## **Anti-CD115 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA6077	Rabbit	H, M, R	WB, IH		
Description	F	Rabbit polyclonal antibody to	0 CD115		
Immunogen	k	KLH-conjugated synthetic pe	otide encompassing a sequence within the center		
	r	region of human CD115. The	exact sequence is proprietary.		
Purification	T	The antibody was purified by	immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of CD115 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/200)		
Gene Symbol	(	CSF1R			
Alternative Na	ames F	FMS; Macrophage colony-sti	mulating factor 1 receptor; CSF-1 receptor; CSF-1-R;		
	(	CSF-1R; M-CSF-R; Proto-onco	gene c-Fms; CD115		
Entrez Gene	1	1436 (Human); 12978 (Mous	e)		
SwissProt	F	P07333 (Human); P09581 (N	ouse); Q00495 (Rat)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	f	freeze/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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kDa A

315

250

180

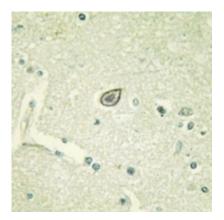
130

95

For research purposes only, not for human use

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Western blot analysis of CD115 expression in SHSY5Y (A) whole cell lysates. (Predicted band size: 107 kD; Observed band size: 175 kD)



Immunohistochemical analysis of CD115 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.

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