

## **Product Data Sheet**

## Anti-CD125 Antibody

Catalog #	Source	Reactivity	Applications		
CPA6205		-			
CPA0205	Rabbit	H, Mk	WB, IH		
Description	I	Rabbit polyclonal antibody t	o CD125		
Immunogen	I	KLH-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	I	region of human CD125. The	exact sequence is proprietary.		
Purification	-	The antibody was purified b	/ immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	ls of CD125 protein.		
Clonality Polyclonal					
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	v	WB (1/500 - 1/1000), IH (1/50	- 1/200)		
Gene Symbol	I	IL5RA			
Alternative Na	ames	IL5R; Interleukin-5 receptor	subunit alpha; IL-5 receptor subunit alpha; IL-5R subunit		
	ä	alpha; IL-5R-alpha; IL-5RA; C	Dw125; CD125		
Entrez Gene	3	3568 (Human)			
SwissProt	(	Q01344 (Human)			
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	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 130

95

72

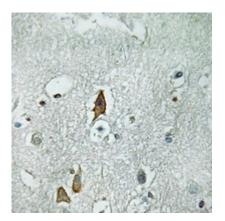
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For research purposes only, not for human use

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Western blot analysis of CD125 expression in A2780 (A), A549 (B) whole cell lysates. (Predicted band size: 47 kD; Observed band size: 75 kD)



Immunohistochemical analysis of CD125 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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