

# **Product Data Sheet**

## **Anti-Cyclophilin F Antibody**

Catalog #	Source	Reactivity	Applications
CPA2454	Rabbit	H, M, R, Rb	WB, IH, IF/IC
Description	F	Rabbit polyclonal antibody to	Cyclophilin F
Immunogen	K	(LH-conjugated synthetic pep	ide encompassing a sequence within the center
	r	egion of human Cyclophilin F.	The exact sequence is proprietary.
Purification	Т	The antibody was purified by i	mmunogen affinity chromatography.
Specificity	F	Recognizes endogenous levels	of Cyclophilin F protein.
Clonality	P	Polyclonal	
Conjugation			
Form	L	iquid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	а	and 0.01% sodium azide.	
Dilution	V	NB (1/500 - 1/1000), IH (1/50 -	1/100), IF/IC (1/50 - 1/200)
Gene Symbol	P	PPIF	
Alternative Na	ames C	CYP3; Peptidyl-prolyl cis-trans	isomerase F mitochondrial; PPIase F; Cyclophilin D;
	C	CyP-D; CypD; Cyclophilin F; Mi	tochondrial cyclophilin; CyP-M; Rotamase F
Entrez Gene	1	10105 (Human); 105675 (Mou	se); 282819 (Rat)
SwissProt	P	230405 (Human); Q99KR7 (Mo	ouse); P29117 (Rat)
Storage/Stabi	ility S	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid
	f	reeze/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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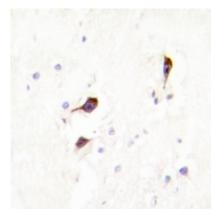
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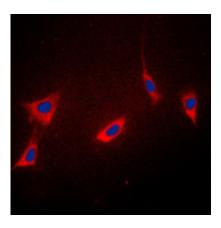
**KDa** A B C D E F 45 35 25 20 15 For research purposes only, not for human use

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Western blot analysis of Cyclophilin F expression in A549 (A), PC12 (B), mouse liver (C), mouse heart (D), rat liver (E), rat heart (F) whole cell lysates. (Predicted band size: 22 kD; Observed band size: 22 kD)



Immunohistochemical analysis of Cyclophilin F 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 Cyclophilin F staining in K562 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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