

# **Product Data Sheet**

### **Anti-OCRL Antibody**

Catalog #	Source	Reactivity	Applications		
CPA1837	Rabbit	H, Mk	WB, IH		
Description	ŀ	Rabbit polyclonal antibody t	o OCRL		
Immunogen	ł	KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
	r	region of human OCRL. The	exact sequence is proprietary.		
Purification	٦	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of OCRL protein.		
Clonality	F	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	١	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	(	OCRL			
Alternative Na	ames l	INPP5F; OCRL1; Inositol poly	phosphate 5-phosphatase OCRL-1; Lowe		
	C	oculocerebrorenal syndrom	e protein		
Entrez Gene		4952 (Human)			
SwissProt	(	Q01968 (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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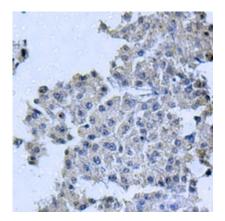
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KDA A B C 180 130 95 72 Western blot analysis of OCRL expression in MG63 (A), A549 (B), HEK293T (C) whole cell lysates. (Predicted band size: 104 kD; Observed band size: 104 kD)



Immunohistochemical analysis of OCRL staining in human liver cancer 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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