

# **Product Data Sheet**

### **Anti-CBR1** Antibody

Catalog #	Source	e Reactivity	Applications		
CPA1167	Rabbit	H, M, R, Mk	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody to	CBR1		
Immunogen		KLH-conjugated synthetic per	otide encompassing a sequence within the center		
		region of human CBR1. The e	exact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous level	s of CBR1 protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/100	) - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol		CBR1			
Alternative N	ames	CBR; CRN; Carbonyl reductas	e [NADPH] 1; 15-hydroxyprostaglandin dehydrogenase		
		[NADP(+)]; NADPH-depender	nt carbonyl reductase 1; Prostaglandin 9-ketoreductase;		
		Prostaglandin-E(2) 9-reducta	se		
Entrez Gene		873 (Human)			
SwissProt		P16152 (Human)			
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
		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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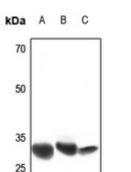
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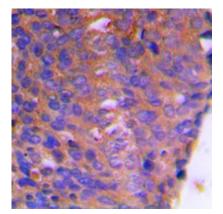
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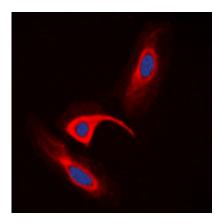
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Western blot analysis of CBR1 expression in mouse liver (A), rat liver (B), rat kidney (C) whole cell lysates. (Predicted band size: 30 kD; Observed band size: 30 kD)



Immunohistochemical analysis of CBR1 staining in human breast 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.



Immunofluorescent analysis of CBR1 staining in HeLa 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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