

Product Data Sheet

Anti-NCOR1 Antibody

Catalog #	Source	Reactivity	Applications	
CPA2409	Rabbit	Н, М	WB, IH, IF/IC	
Description	Rat	bit polyclonal antibody	to NCOR1	
Immunogen	KLF	I-conjugated synthetic	peptide encompassing a sequence within the N-term	
	reg	ion of human NCOR1. T	he exact sequence is proprietary.	
Purification	The	e antibody was purified	by immunogen affinity chromatography.	
Specificity	Rec	cognizes endogenous le	vels of NCOR1 protein.	
Clonality	Pol	yclonal		
Conjugation				
Form	Liqu	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
	and	d 0.01% sodium azide.		
Dilution	WB	s (1/500 - 1/1000), IH (1/2	.00 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	NC	OR1		
Alternative Na	ames KIA	A1047; Nuclear recepto	or corepressor 1; N-CoR; N-CoR1	
Entrez Gene	961	L1 (Human)		
SwissProt	07	O75376 (Human); Q60974 (Mouse)		
Storage/Stabi	lity Shi	pped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid	
	free	eze/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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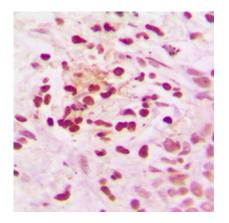
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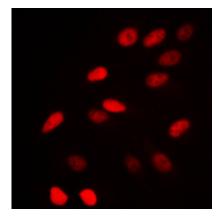
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Western blot analysis of NCOR1 expression in HepG2 (A), CT26 (B) whole cell lysates. (Predicted band size: 270 kD; Observed band size: 320 kD)



Immunohistochemical analysis of NCOR1 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 NCOR1 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.

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