

## **Product Data Sheet**

### **Anti-NDRG2** Antibody

Catalog #	Source	Reactivity	Applications		
CPA4113	Rabbit	H, M, R, B, P	WB, IH		
Description	Ra	abbit polyclonal antibody to	NDRG2		
Immunogen	KI	LH-conjugated synthetic pep	tide encompassing a sequence within the center		
	re	egion of human NDRG2. The	exact sequence is proprietary.		
Purification	Tł	he antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous level	s of NDRG2 protein.		
Clonality	Ро	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	VB (1⁄500 - 1⁄1000), IH (1⁄100	- 1/200)		
Gene Symbol	Ν	IDRG2			
Alternative Na	ames Kl	IAA1248; SYLD; Protein NDR	G2; N-myc downstream-regulated gene 2 protein;		
	Рі	rotein Syld709613			
Entrez Gene	57	7447 (Human); 29811 (Mou	se); 171114 (Rat)		
SwissProt	Q	Q9UN36 (Human); Q9QYG0 (I	Mouse); Q8VBU2 (Rat)		
Storage/Stabi	<b>lity</b> Sł	hipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	fr	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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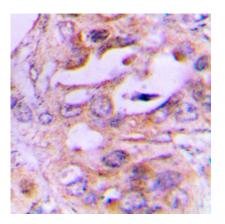
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kDa A B C

S S Western bl For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of NDRG2 expression in HeLa (A), Raw264.7 (B), PC12 (C) whole cell lysates. (Predicted band size: 40 kD; Observed band size: 40 kD)



Immunohistochemical analysis of NDRG2 staining in human lung 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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