

# **Product Data Sheet**

### **Anti-LYN Antibody**

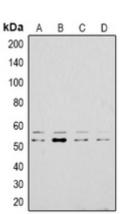
Catalog #	Source	Reactivity	Applications	
CPA4957	Rabbit	н, м	WB, IH	
Description		Rabbit polyclonal antibody	to LYN	
Immunogen		KLH-conjugated synthetic pe	eptide encompassing a sequence within the C-term	
		region of human LYN. The e	kact sequence is proprietary.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of LYN 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 (1/50	) - 1/200)	
Gene Symbol		LYN		
Alternative Na	ames	JTK8; Tyrosine-protein kinas	e Lyn; Lck/Yes-related novel protein tyrosine kinase;	
		V-yes-1 Yamaguchi sarcoma	viral related oncogene homolog; p53Lyn; p56Lyn	
Entrez Gene		4067 (Human); 17096 (Mouse)		
SwissProt		P07948 (Human); P25911 (N	/louse)	
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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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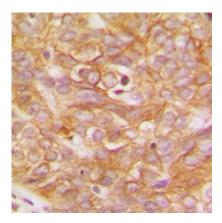




For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of LYN expression in A431 (A), Hela (B), mouse kidney (C), PC12 (D) whole cell lysates. (Predicted band size: 58 kD; Observed band size: 53; 56 kD)



Immunohistochemical analysis of LYN 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.

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