

# **Product Data Sheet**

# Anti-CD43 Antibody

Catalog #	Source	Reactivity	Applications	
CPA9606	Mouse	Н	IH	
Description		Mouse monoclonal antibod	/ to CD43	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within human CD43.	
		The exact sequence is propr	ietary.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of CD43 protein.	
Clonality		Monoclonal		
Conjugation				
Form		Mouse IgG1. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium		
	;	azide.		
Dilution		IH (1/100 - 1/300)		
Gene Symbol	:	SPN		
Alternative Na	ames	CD43; Leukosialin; Galactog	ycoprotein; GALGP; Leukocyte sialoglycoprotein;	
	:	Sialophorin; CD43		
Entrez Gene		101929889, 6693 (Human)		
SwissProt		P16150 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	t	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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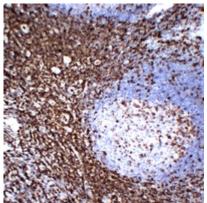
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# Coherion

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Immunohistochemical analysis of CD43 staining in human anaplastic large cell lymphoma 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 t



Immunohistochemical analysis of CD43 staining in human tonsil 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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