

Product Data Sheet

Anti-CD104 Antibody

Catalas #	Courses	Department		Applications
Catalog #	Source	Reactivity	y	Applications
CPA1619	Rabbit	Н <i>,</i> М		WB, IH
Description	Ra	abbit polyclonal	antibody to CD104	
Immunogen	KL	LH-conjugated sy	nthetic peptide encompa	assing a sequence within the C-term
	re	egion of human C	CD104. The exact sequence	ce is proprietary.
Purification	Th	ne antibody was	purified by immunogen a	affinity chromatography.
Specificity	Re	ecognizes endog	enous levels of CD104 pro	otein.
Clonality	Pc	olyclonal		
Conjugation				
Form	Lic	quid in 0.42% Pc	tassium phosphate, 0.87	% Sodium chloride, pH 7.3, 30% glycerol,
	an	nd 0.01% sodium	n azide.	
Dilution	W	/B (1/500 - 1/1000	0), IH (1/100 - 1/200)	
Gene Symbol	IT	GB4		
Alternative Na	ames Int	tegrin beta-4; G	P150; CD104	
Entrez Gene	36	691 (Human); 19	2897 (Mouse)	
SwissProt	P1	16144 (Human);	A2A863 (Mouse)	
Storage/Stabi	lity Sh	nipped at 4 $^\circ~$ C. (Jpon delivery aliquot and	l store at -20 $^\circ~$ C for one year. Avoid
	fre	eeze/thaw cycles	S.	

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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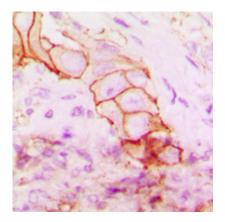
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Western blot analysis of CD104 expression in A431 (A), SW480 (B) whole cell lysates. (Predicted band size: 106; 194; 195; 200; 202 kD; Observed band size: 202 kD)



Immunohistochemical analysis of CD104 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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