

Product Data Sheet

Anti-BAF155 Antibody

Catalog #	Source	e Reactivity	Applications	
CQA1296	Rabbit	: Н, R	WB, IH, IP, ChIP	
Description		Rabbit polyclonal antibody t	o BAF155	
Immunogen		Recombinant full length pro	tein of human BAF155	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of BAF155 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 - 1/2000), IH (1/50 - 1/200), IP (1/50 - 1/200), ChIP (1/20 - 1/100)		
Gene Symbol		SMARCC1		
Alternative Names		BAF155; SWI/SNF complex subunit SMARCC1; BRG1-associated factor 155; BAF155;		
		SWI/SNF complex 155 kDa s	ubunit; SWI/SNF-related matrix-associated	
		actin-dependent regulator of	f chromatin subfamily C member 1	
Entrez Gene		6599 (Human)		
SwissProt		Q92922 (Human)		
Storage/Stabi	lity	Shipped at 4 $^\circ$ C. Upon deliv	ery aliquot and store at -20 $^\circ$ 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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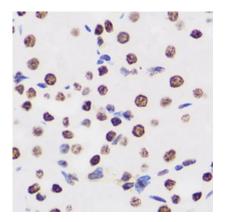
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For research purposes only, not for human use

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Western blot analysis of BAF155 expression in SW620 (A), HEK293T (B) whole cell lysates. (Predicted band size: 122 kD; Observed band size: 155 kD)



Immunohistochemical analysis of BAF155 staining in human kidney 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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