

Product Data Sheet

Anti-INI1 Antibody

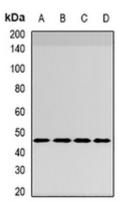
Catalog #	Source	e Reactivity	Applications	
CQA2242	Rabbit	Н, М, R	WB, IH, IF/IC, IP, ChIP	
Description		Rabbit polyclonal antibody t	o INI1	
Immunogen		Recombinant full length pro	tein of human INI1	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of INI1 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium p	hosphate, 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), IF/IC (1/50 - 1/200), IP (1/50 - 1/200), ChIP (1/20 -	
		1/100)		
Gene Symbol		SMARCB1		
Alternative Na	ames	BAF47; INI1; SNF5L1; SWI/SI	NF-related matrix-associated actin-dependent regulator	
		of chromatin subfamily B me	ember 1; BRG1-associated factor 47; BAF47; Integrase	
		interactor 1 protein; SNF5 h	omolog; hSNF5	
Entrez Gene		6598 (Human); 20587 (Mouse)		
SwissProt		Q12824 (Human); Q9Z0H3 (Mouse)	
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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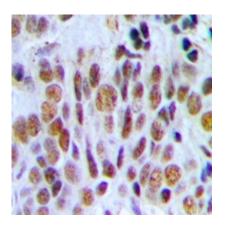




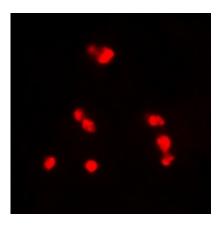
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Western blot analysis of INI1 expression in HepG2 (A), MCF7 (B), mouse brain (C), mouse lung (D) whole cell lysates. (Predicted band size: 43; 44 kD; Observed band size: 44 kD)



Immunohistochemical analysis of INI1 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.



Immunofluorescent analysis of INI1 staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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