

Product Data Sheet

Anti-HEY1 Antibody

Catalog #	Source	e Reactivity	Applications		
CQA4671	Rabbit	Н, М	WB, IF/IC		
Description		Rabbit polyclonal antibody t	o HEY1		
Immunogen		KLH-conjugated synthetic peptide of human HEY1. The exact sequence is			
		proprietary.			
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of HEY1 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), IF/IC (1/	50 - 1/200)		
Gene Symbol		HEY1			
Alternative Names		BHLHB31; CHF2; HERP2; HESR1; HRT1; Hairy/enhancer-of-split related with YRPW			
		motif protein 1; Cardiovascu	lar helix-loop-helix factor 2; CHF-2; Class B basic		
		helix-loop-helix protein 31; l	HLHb31; HES-related repressor protein 1; Hairy and		
		enhancer of split-related pro	tein 1; HESR-1; Hairy-related transcription factor 1;		
		HRT-1; hHRT1			
Entrez Gene		23462 (Human)			
SwissProt		Q9Y5J3 (Human); Q9WV93 (Mouse)		
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

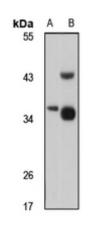
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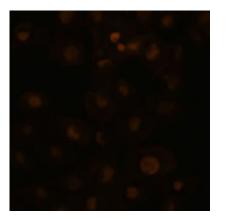
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Western blot analysis of HEY1 expression in HEK293T (A), mouse heart (B) whole cell lysates. (Predicted band size: 32; 33 kD; Observed band size: 33 kD)



Immunofluorescent analysis of HEY1 staining in A431 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 $^{\circ}$ C in a humidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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