

Product Data Sheet

Anti-IL-23R Antibody

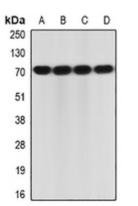
Catalog #	Source	Reactivity	Applications
CQA2033	Rabbit	Н, М	WB, IH, IF/IC
Description		Rabbit polyclonal antibody	to IL-23R
Immunogen		Recombinant full length pro	otein of human IL-23R
Purification		The antibody was purified l	by immunogen affinity chromatography.
Specificity		Recognizes endogenous lev	els of IL-23R 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 (1/5	0 - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol		IL23R	
Alternative Na	ames	Interleukin-23 receptor; IL-	23 receptor; IL-23R
Entrez Gene		149233 (Human); 209590 (Mouse)
SwissProt		Q5VWK5 (Human); Q8K4B4	ł (Mouse)
Storage/Stabi	lity	Shipped at 4 $^{\circ}$ C. Upon deli	very 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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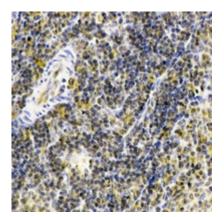




For research purposes only, not for human use

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Western blot analysis of IL-23R expression in K562 (A), HepG2 (B), mouse intestine (C), mouse spleen (D) whole cell lysates. (Predicted band size: 20; 25; 30; 40; 42; 71 kD; Observed band size: 71 kD)



Immunohistochemical analysis of IL-23R staining in human spleen 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 IL-23R staining in HepG2 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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