

Product Data Sheet

Anti-IL-25 Antibody

Catalog #	Source	e Reactivity	Applications
CQA3700	Rabbit	Н	WB, IH, IF/IC
Description		Rabbit polyclonal antibod	y to IL-25
Immunogen		Recombinant protein of h	uman IL-25. The exact sequence is proprietary.
Purification		The antibody was purified	l by immunogen affinity chromatography.
Specificity		Recognizes endogenous le	evels of IL-25 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 - 1/1000), IH (1,	/100 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol		IL25	
Alternative Na	ames	IL17E; Interleukin-25; IL-2	5; Interleukin-17E; IL-17E
Entrez Gene		64806 (Human)	
SwissProt		Q9H293 (Human)	
Storage/Stabi	lity	Shipped at 4 $^{\circ}$ C. Upon de	livery 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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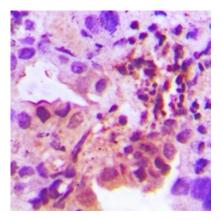
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Western blot analysis of IL-25 expression in U937 (A) whole cell lysates. (Predicted band size: 20 kD; Observed band size: 25 kD)



Immunohistochemical analysis of IL-25 staining in human prostate 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-25 staining in U937 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 hidified chamber. Cells were washed with PBST and incubated with a Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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