

## **Product Data Sheet**

## Anti-IL-9 Antibody

Catalog #	Source	Reac	tivity	Applicat	ions	
CQA3699	Rabbit	H, M,	, R	WB, IH		
Description	I	Rabbit polyclo	onal antibody to I	L-9		
Immunogen		Recombinant protein of human IL-9. The exact sequence is proprietary.				
Purification	-	The antibody	tibody was purified by immunogen affinity chromatography.			
Specificity	ļ	Recognizes endogenous levels of IL-9 protein.				
Clonality	ļ	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	i	and 0.01% soc	dium azide.			
Dilution	,	WB (1/500 - 1/	1000), IH (1/100 -	1/200)		
Gene Symbol	I	IL9				
Alternative Names		Interleukin-9; IL-9; Cytokine P40; T-cell growth factor P40				
Entrez Gene 357		3578 (Human); 16198 (Mouse)				
SwissProt	I	P15248 (Human); P15247 (Mouse)				
Storage/Stabi	lity :	Shipped at 4 $^\circ$	C. Upon deliver	y aliquot and store at -	$20^\circ$ C for one year. Avoid	
	t	freeze/thaw c	ycles.			

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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**kDa** 70

> 51 38 28

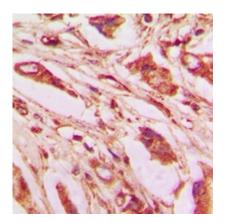
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For research purposes only, not for human use

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Western blot analysis of IL-9 expression in human spleen (A) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 15 kD)



Immunohistochemical analysis of IL-9 staining in human lung 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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