

## **Product Data Sheet**

### Recombinant Anti-Cytokeratin 18 Rabbit mAb

Catalog #	Source	Reactivity	Applications			
CMA1005	Rabbit	H, M, R	WB, IH			
Description	Re	Recombinant rabbit monoclonal antibody to Cytokeratin 18				
Immunogen	Re	ecombinant fusion protein	of human Cytokeratin 18. The exact sequence is			
	рг	roprietary.				
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.			
Specificity	Re	ecognizes endogenous leve	els of Cytokeratin 18 protein.			
Clonality	Μ	Ionoclonal				
Conjugation						
Form	Li	quid in PBS, pH 7.3, 50% g	ycerol, and 0.05% Proclin300.			
Dilution	Ŵ	/B (1/500 - 1/1000), IH (1/10	0 - 1/200)			
Gene Symbol	KI	RT18				
Alternative Na	ames C	YK18; Keratin, type I cytosł	eletal 18; Cell proliferation-inducing gene 46 protein;			
	Cy	ytokeratin-18; CK-18; Kerat	in-18; K18			
Entrez Gene	38	875 (Human); 16668 (Mou	se); 294853 (Rat)			
SwissProt	P	05783 (Human); P05784 (N	1ouse); Q5BJY9 (Rat)			
Storage/Stabi	<b>lity</b> Sł	nipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	fr	eeze/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

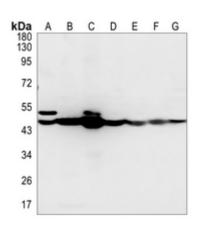
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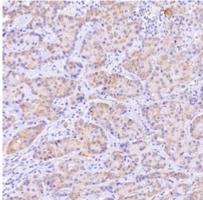
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Western blot analysis of Cytokeratin 18 expression in HEK293T (A), EC9706 (B), SGC7901 (C), mouse liver (D), mouse kidney (E), rat liver (F), rat kidney (G) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 48 kD)



Immunohistochemical analysis of Cytokeratin 18 staining in human liver 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.

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