

Product Data Sheet

Recombinant Anti-ASS1 Rabbit mAb

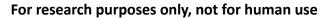
Catalog #	Source	Reactivity	Applications
CMA4595	Rabbit	H, M, R	WB, IH
Description	Rec	ombinant rabbit mono	clonal antibody to ASS1
Immunogen	KLF	I-conjugated synthetic	peptide encompassing a sequence within human ASS1
	pro	tein. The exact sequen	ce is proprietary.
Purification	The	antibody was purified	by immunogen affinity chromatography.
Specificity	Rec	ognizes endogenous le	vels of ASS1 protein
Clonality	Мо	noclonal	
Conjugation			
Form	Liqu	uid in PBS containing 50	0% glycerol, 0.2% BSA and 0.01% sodium azide.
Dilution	WB	(1/500 - 1/1000), IH (1/5	50 - 1/200)
Gene Symbol	ASS	51	
Alternative Na	ames ASS	; Argininosuccinate syr	thase; Citrullineaspartate ligase
Entrez Gene	445	i (Human); 11898 (Μοι	ise); 25698 (Rat)
SwissProt	POC	966 (Human); P16460	(Mouse); P09034 (Rat)
Storage/Stabi	lity Shij	oped at 4 $^\circ$ C. Upon del	ivery aliquot and store at -20 $^\circ~$ C for one year. Avoid
	free	eze/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

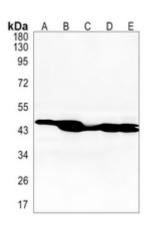
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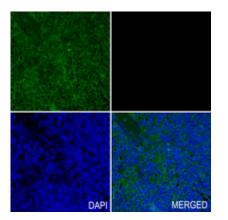




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Western blot analysis of ASS1 expression in MCF7 (A), mouse liver (B), mouse kidney (C), rat liver (D), rat kidney (E) whole cell lysates. (Predicted band size: 46 kD; Observed band size: 47 kD)



Immunohistochemical analysis of ASS1 staining in human tonsil 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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