

## **Product Data Sheet**

### **Anti-MINA53 Antibody**

Catalog #	Source	Reactivity	Applications			
CPA3086	Rabbit	H, M, R	WB, IF/IC			
Description	Ra	Rabbit polyclonal antibody to MINA53				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center				
	re	gion of human MINA53. T	ne exact sequence is proprietary.			
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.			
Specificity	Re	ecognizes endogenous leve	ls of MINA53 protein.			
Clonality	Pc	blyclonal				
Conjugation						
Form	Lie	quid in 0.42% Potassium p	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.				
Dilution	W	′B (1/500 - 1/1000), IF/IC (1/	50 - 1/200)			
Gene Symbol	М	INA				
Alternative Na	ames M	DIG; MINA53; NO52; Bifur	ctional lysine-specific demethylase and			
	hi	stidyl-hydroxylase MINA; 6	OS ribosomal protein L27a histidine hydroxylase;			
	Hi	istone lysine demethylase	MINA; MYC-induced nuclear antigen; Mineral			
	dı	ust-induced gene protein; I	Nucleolar protein 52; Ribosomal oxygenase MINA; ROX			
Entrez Gene	84	1864 (Human); 67014 (Mo	use); 266670 (Rat)			
SwissProt	Q	8IUF8 (Human); Q8CD15 (I	Mouse); Q8CFC1 (Rat)			
Storage/Stabi	<b>lity</b> Sh	nipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid			
	fre	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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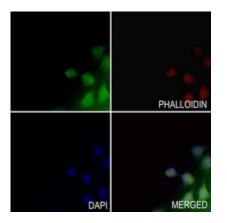
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Western blot analysis of MINA53 expression in rat liver (A) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 53 kD)



Immunofluorescent analysis of MINA53 staining in A549 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 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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