

Product Data Sheet

Anti-NFS1 Antibody

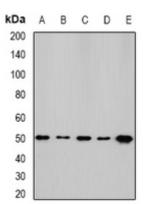
Catalog #	Source	Reactivity	Applications		
CQA1440	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	bbit polyclonal antibody	/ to NFS1		
Immunogen	Re	combinant full length pi	rotein of human NFS1		
Purification	Th	e antibody was purified	by immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous le	vels of NFS1 protein.		
Clonality	Ро	lyclonal			
Conjugation					
Form	Lic	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/2000), IH (1/5	50 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	NF	S1			
Alternative Na	ames NI	FS; Cysteine desulfurase	mitochondrial		
Entrez Gene	90	54 (Human); 18041 (Mc	puse)		
SwissProt	QS	9Y697 (Human); Q9Z1J3	(Mouse); Q99P39 (Rat)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon deliv	very 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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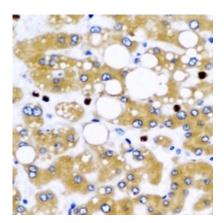




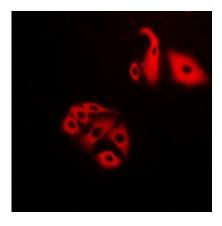
For research purposes only, not for human use

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Western blot analysis of NFS1 expression in HepG2 (A), SKOV3 (B), mouse kidney (C), mouse heart (D), rat liver (E) whole cell lysates. (Predicted band size: 44; 50 kD; Observed band size: 47 kD)



Immunohistochemical analysis of NFS1 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.



Immunofluorescent analysis of NFS1 staining in Hela 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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