

# **Product Data Sheet**

## Anti-FDFT1 Antibody

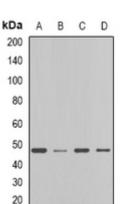
Catalog #	Source	Reactivity	Applications
CQA1309	Rabbit	H, M, R	WB, IH, IF/IC
Description	Ra	bbit polyclonal antibody	to FDFT1
Immunogen	Re	combinant full length pro	otein of human FDFT1
Purification	Th	e antibody was purified b	by immunogen affinity chromatography.
Specificity	Re	cognizes endogenous lev	els of FDFT1 protein.
Clonality	Ро	lyclonal	
Conjugation			
Form	Lic	quid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	d 0.01% sodium azide.	
Dilution	W	B (1/500 - 1/2000), IH (1/50	) - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	FD	FT1	
Alternative Na	ames Sq	ualene synthase; SQS; SS	; FPP:FPP farnesyltransferase; Farnesyl-diphosphate
	far	rnesyltransferase	
Entrez Gene	22	22 (Human); 14137 (Mou	ise); 29580 (Rat)
SwissProt	Р3	7268 (Human); P53798 (I	Mouse); Q02769 (Rat)
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry 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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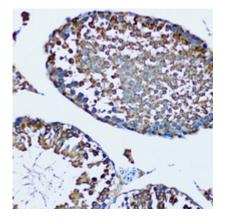




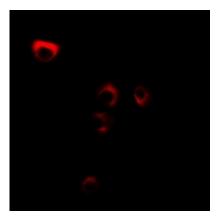
For research purposes only, not for human use

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Western blot analysis of FDFT1 expression in MCF7 (A), HT29 (B), mouse liver (C), rat liver (D) whole cell lysates. (Predicted band size: 35; 38; 40; 43; 48 kD; Observed band size: 48 kD)



Immunohistochemical analysis of FDFT1 staining in mouse testis 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 FDFT1 staining in MCF7 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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