

Anti-HADHA Antibody (A14688)

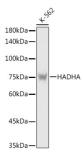
Specifications:

Name:	Anti-HADHA Antibody
Description:	Rabbit polyclonal antibody to HADHA.
Applications:	WB, ICC/IF, IP
Recommended Dilutions:	WB: 1:2,000-1:6,000, ICC/IF: 1:50-1:200, IP: 1:50-1:100
Reactivity:	Human, Mouse, Rat
Immunogen:	Recombinant fusion protein containing a sequence corresponding to amino acids 484-763 of human HADHA (NP_000173.2).
Sequence:	IAAVSKRPEKVIGMHYFSPVDKMQLLEIITTEKTSKDTSASAVAVGLKQGKVIIVVKD GPGFYTTRCLAPMMSEVIRILQEGVDPKKLDSLTTSFGFPVGAATLVDEVGVDVAKHV AEDLGKVFGERFGGGNPELLTQMVSKGFLGRKSGKGFYIYQEGVKRKDLNSDMDSILA SLKLPPKSEVSSDEDIQFRLVTRFVNEAVMCLQEGILATPAEGDIGAVFGLGFPPCLG GPFRFVDLYGAQKIVDRLKKYEAAYGKQFTPCQLLADHANSPNKKFYQ
Host:	Rabbit
Clonality:	Polyclonal
Isotype:	lgG
Conjugate:	Uncertaged
	Unconjugated
Purification:	Affinity purification.
Purification: Molecular Weight:	
	Affinity purification.
Molecular Weight:	Affinity purification. 78 kDa
Molecular Weight: Product Form:	Affinity purification. 78 kDa Liquid Supplied in Phosphate Buffered Saline, pH 7.3, with 50% Glycerol and 0.02% Sodium

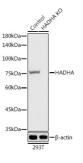
antibodies

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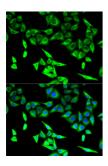
Images:



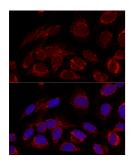
Western blot analysis of extracts of K-562 cells, using Anti-HADHA Antibody (A14688) at 1:5,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 1s.



Western blot analysis of extracts from normal (control) and HADHA knockout (KO) 293T cells, using Anti-HADHA Antibody (A14688) at 1:5,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 1s.



Immunofluorescence analysis of U2OS cells using Anti-HADHA Antibody (A14688). DAPI was used to stain the cell nuclei (blue).



Confocal immunofluorescence analysis of U-2 OS cells using Anti-HADHA Antibody (A14688) at a dilution of 1:100. DAPI was used to stain the cell nuclei (blue).